



ATRIAL FIBRILLATION

Radhika Nandur Bukkapatnam MD MAAS FACC

11/30/23

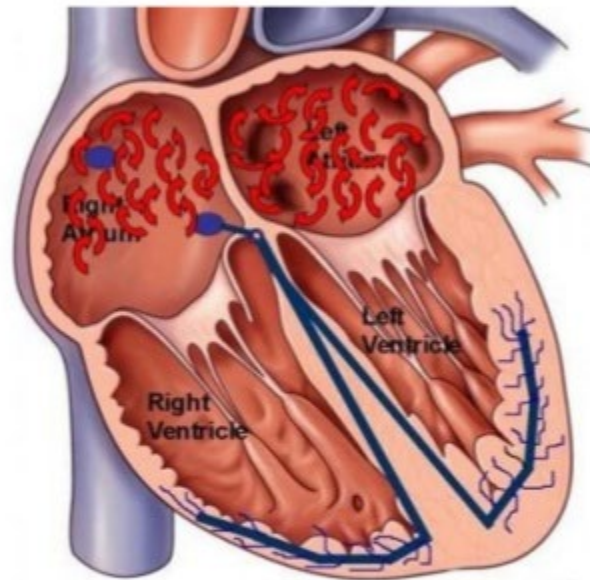
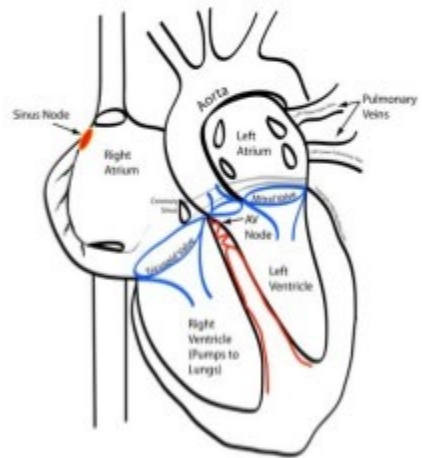


OBJECTIVES

- Introduction
- Classification
- Burden of the problem
- Diagnosis
- Management

DEFINITION

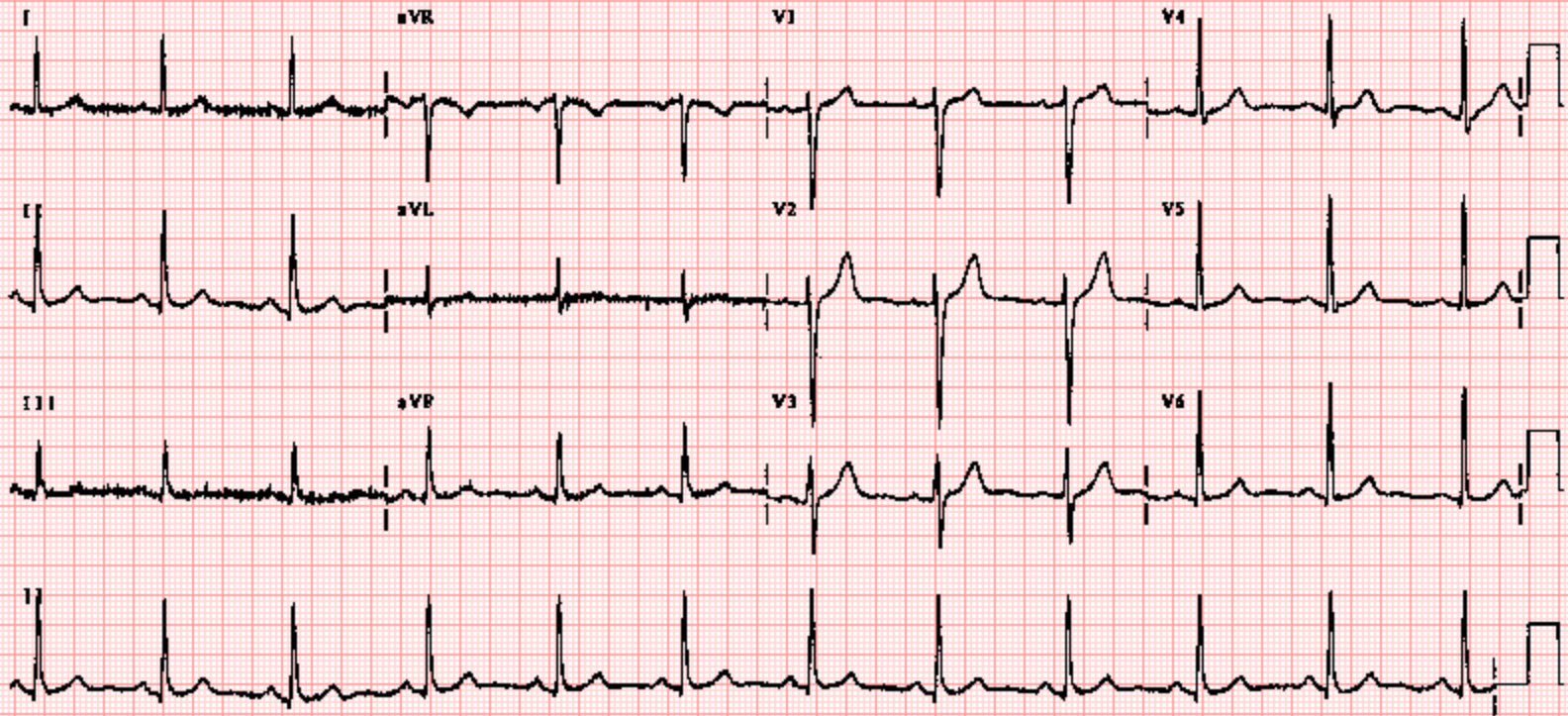
- Irregular, chaotic heart rhythm that comes from the left atrium or the pulmonary veins.
- Paroxysmal
- Persistent
- Chronic



DIAGNOSIS

- Pulse palpation
- 12 lead ECG
- Holter monitoring
- **Others**
- Echocardiogram, CXR
- TFT, Electrolytes, Clotting, LFT,CBC

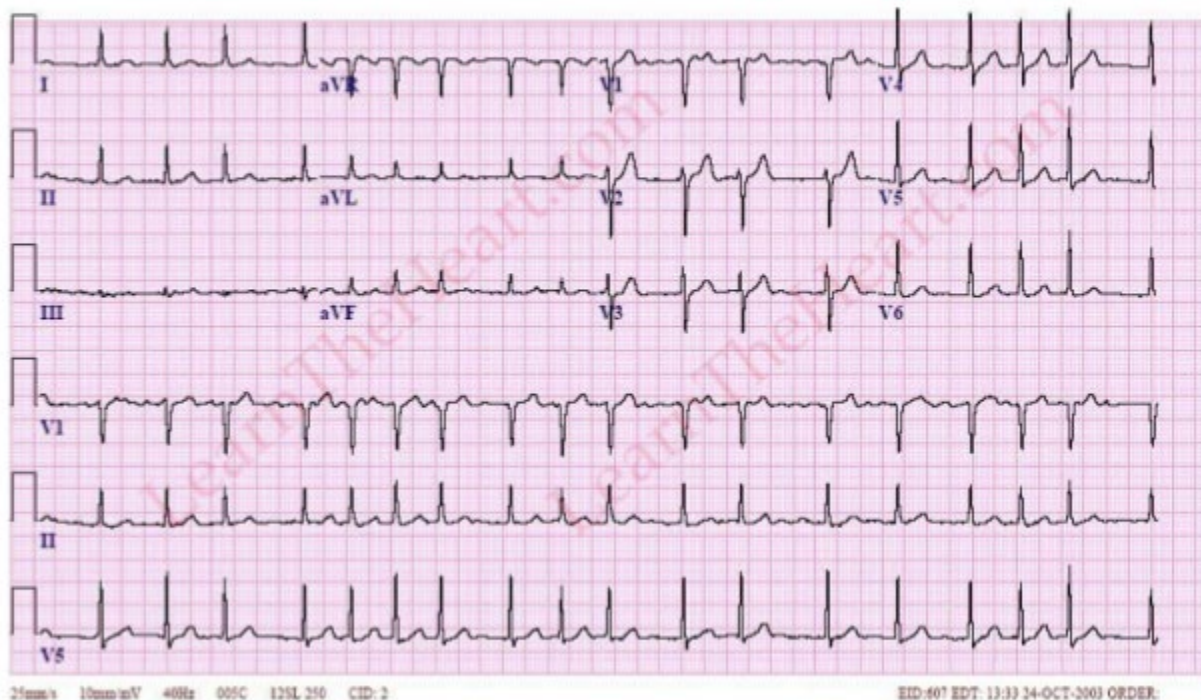
Normal ECG



I.CC 0000-0000 Speed: 25 mm/sec Limb: 10 mV Chest: 10 mm/mV

50% 0.15-150 Hz

16405



Prevalance

- 2.2 Million people in the US
- 6.5 cases/1000 examinations
- 4% > 60yrs
- 8 % > 80 yrs
- 25% of individuals aged 40 yrs and older will develop AF in their life time.

Clinical events (outcomes) affected by AF

Outcome Parameter	Relative change in AF patients
1. Death	1. Death rate is doubled
2. Stroke	2. Stroke risk increases 5 times
3. Hospitalisation	3. More frequent
4. Quality of life and exercise capacity	4. Can be markedly decreased
5. LV function	5. Tachycardiomyopathy/ heart failure

Classification of AF

Terminology	Clinical features	
Initial event (first detected episode)	Symptomatic Asymptomatic Onset unknown	Rhythm/Rate
Paroxysmal	Spontaneous termination <7 days and most often <48 hours	Rhythm Control
Persistent	Not self-terminating Lasting >7 days or prior cardioversion	Rhythm or Rate control
Permanent (‘accepted’)	Not terminated Terminated but relapsed No cardioversion attempt	Rate Control

Etiologies of AF

CARDIAC

Hypertensive heart disease

Valvular heart disease

Ischaemic heart disease

Cardiomyopathy

Pericarditis

Congenital heart disease

Post Cardiac surgery

Etiologies of AF contd:

NON CARDIAC

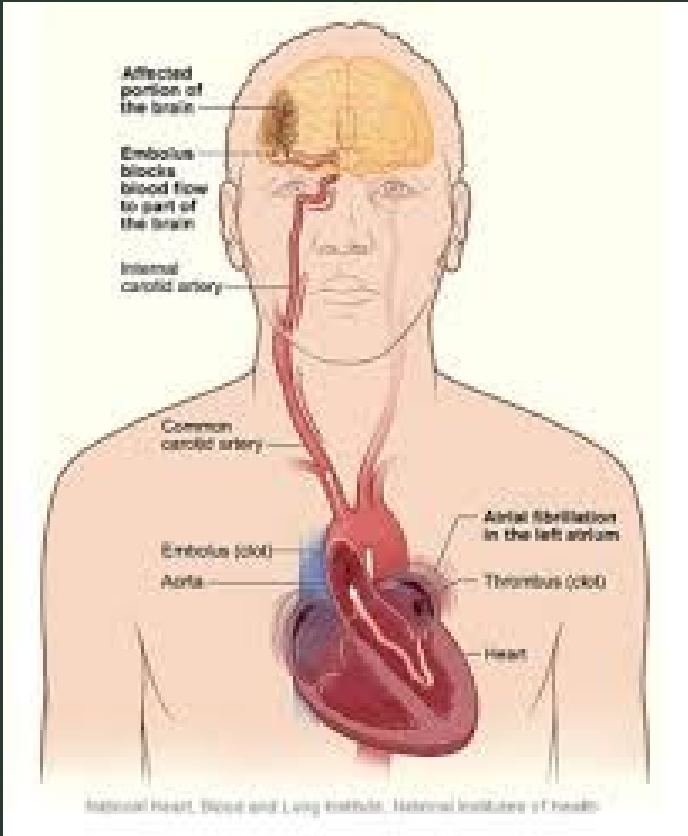
1. Pulmonary : Pneumonia, COPD,PE
2. Hyperthyroidism
3. Excess catecholamine /sympathetic activity
4. Drugs and alcohol
5. Significant electrolyte imbalance

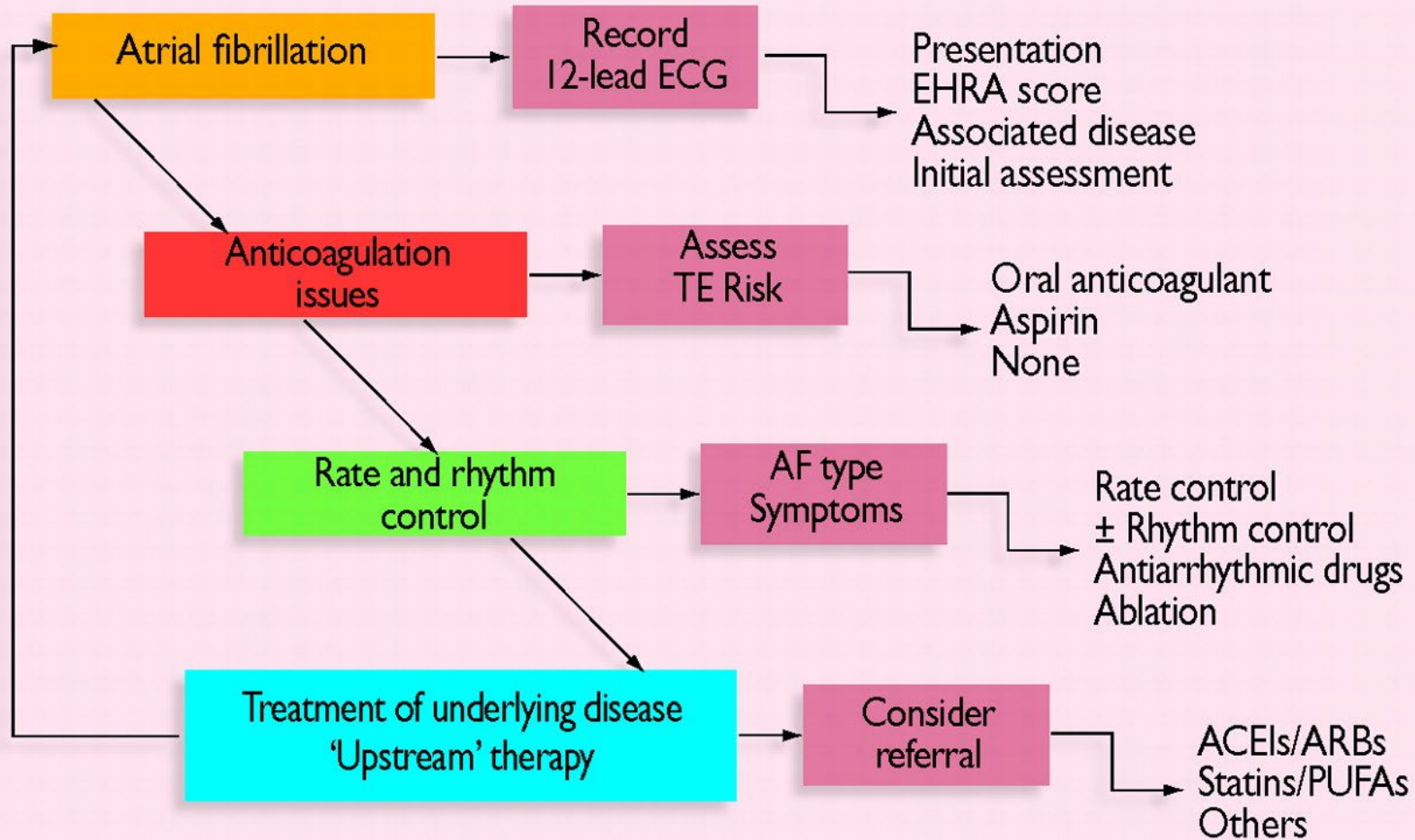
LONE ATRIAL FIBRILLATION

- Younger patients < 60
- No underlying cause
- Usually not much symptoms
- Normal heart structure
- No associated co-morbidities

Why AF management is important?

- extremely common
- Can lead to symptoms
- potentially serious consequences:
 - embolism
 - impaired cardiac output
 - increased mortality





MANAGEMENT OF A FIB

- Acute Management
- Stroke Prevention
- Rate control: For symptoms and to prevent decompensation
- Rhythm control: To prevent stroke, dementia, CHF
- Prevention



ACUTE TREATMENT OF A FIB

Management of Acute AF (<48 hrs)

- Haemodynamically unstable :
hypotension/heart failure/chest pain/syncope

Use DC Cardioversion

Haemodynamically stable :

Rate control : If significant tachycardia

Rhythm control : Flecainide, Propafenone (cl-I)
Amiodarone, Sotalol (cl-III)

Anticoagulant : LMWH



STROKE PREVENTION



How do we determine stroke risk ?

- **0 points – low risk** (1.2-3.0 strokes per 100 patient years)
- **1-2 points – moderate risk** (2.8-4.0 strokes per 100 patient years)
- **≥ 3 points – high risk** (5.9-18.2 strokes per 100 patient years)

Stroke Prevention

CHADSVasc Score

- C – congestive heart failure
 - H – hypertension
 - A – age 65 (1 pt), age 75 (2 pts)
 - D – diabetes
 - S – prior stroke or TIA
 - Va - vascular disease
 - S – gender (female gender – now no longer considered)
-
- CHADsVasc ≥ 2 – advise anticoagulation
 - CHADSVasc score ≥ 1 – consider anticoagulation
 - CHADSVasc score 0 – anticoagulation not advised

Annualized Stroke Risk vs CHADS-VASc Score in Afib Patients

CHADS2 – VASc Score	
C	Congestive Heart Failure
H	Hypertension (>140/90 mmHg)
A	Age ≥ 75
D	Diabetes Mellitus
S₂	Prior TIA or stroke
V	Vascular disease (MI, aortic plaque etc)
A	Age 65-74
Sc	Sex category (Female = 1 pt)

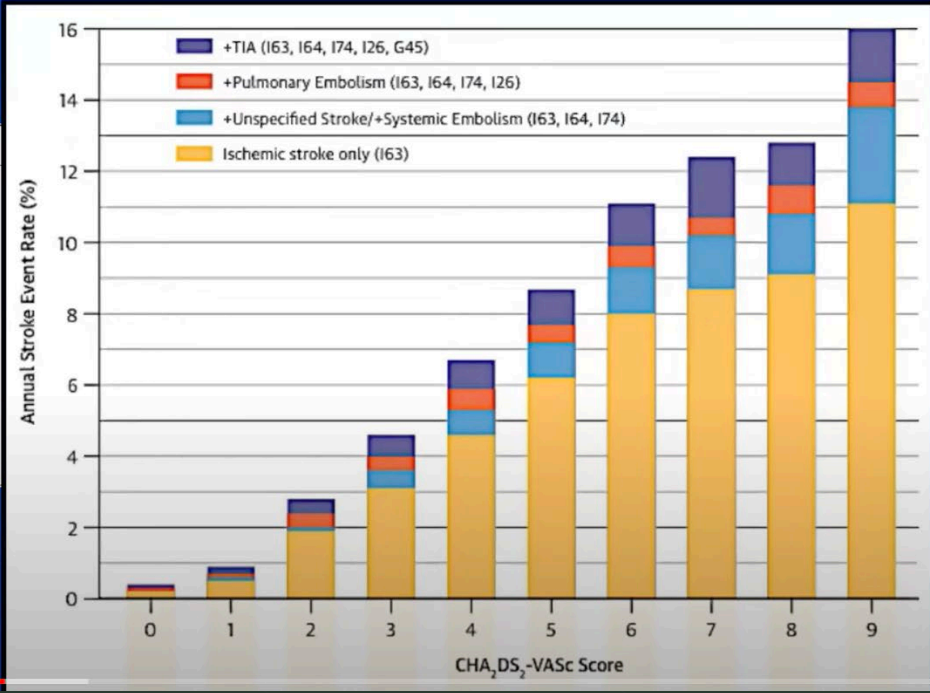


Table 1. Changes in exposure to oral anticoagulants, 2015 Q4 to 2016 Q4*

Drug Name	Dispensed Prescriptions		Percent Change	Market Share**
	2015 Q4	2016 Q4		
apixaban	1,315,213	2,183,821	66.0%	19.2%
dabigatran etexilate	487,527	486,176	-0.3%	4.3%
edoxaban	23,563	23,886	1.4%	0.2%
rivaroxaban	1,948,201	2,209,216	13.4%	19.4%
warfarin	7,332,251	6,488,962	-11.5%	57.0%
Total oral anticoagulants	11,106,755	11,392,061	2.6%	

* Dispensed outpatient prescriptions per QuintilesIMS data

** In 2016 Q4

Types of Blood Thinners

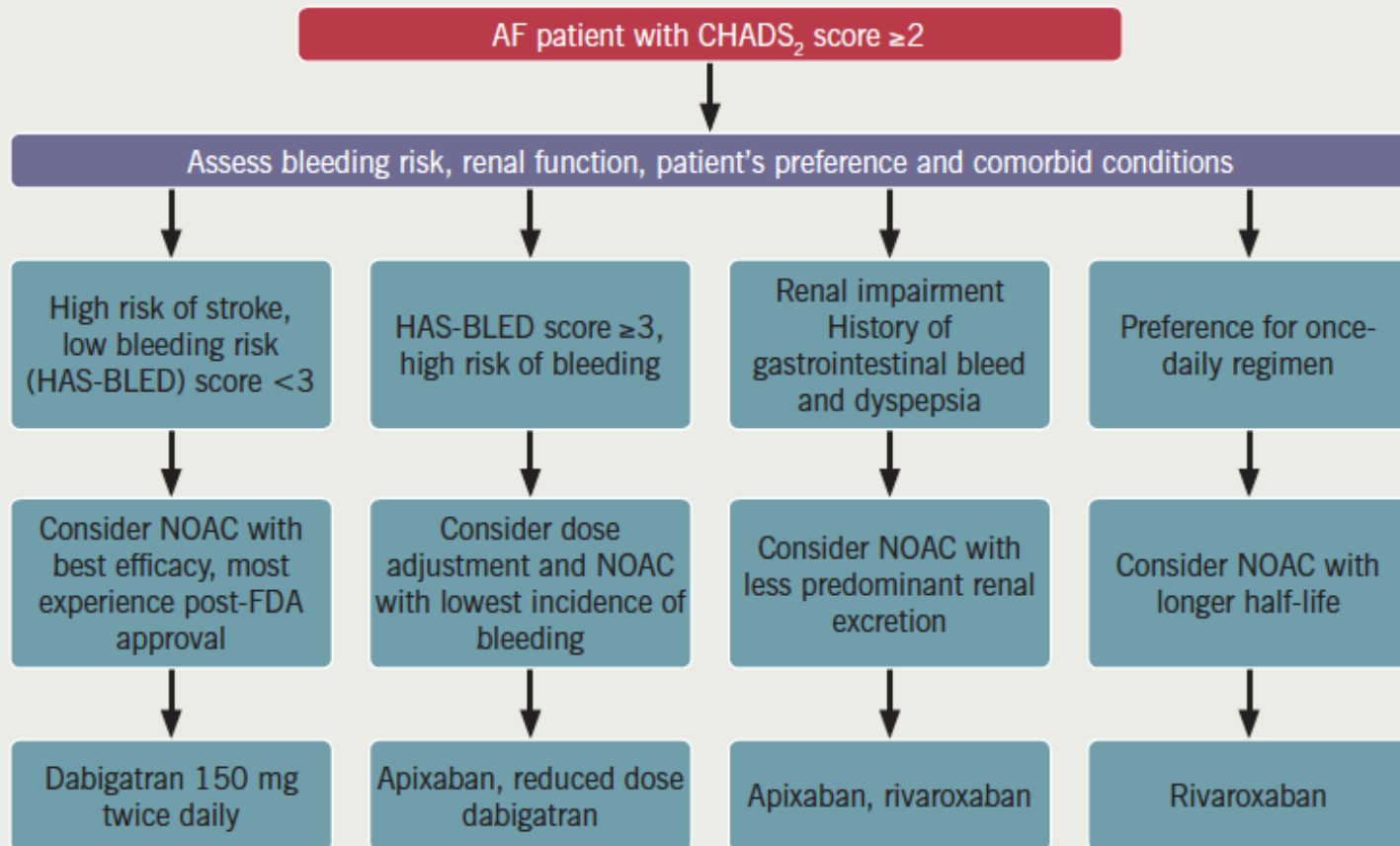
The HAS-BLED bleeding risk score

Letter	Clinical characteristic*	Points awarded
H	Hypertension	1
A	Abnormal renal and liver function (1 point each)	1 or 2
S	Stroke	1
B	Bleeding	1
L	Labile INRs	1
E	Elderly (e.g. age > 65 years)	1
D	Drugs or alcohol (1 point each)	1 or 2
		Maximum 9 points

Bleeding Risk

- Assessment of bleeding risk should be part of the clinical assessment of AF patients prior to starting anticoagulation
- Antithrombotic benefits and potential bleeding risks of long-term coagulation should be explained and discussed with the patient
- Aim for a target INR of between 2.0 and 3.0
- Forms of monitoring include point of care or near patient testing and patient self-monitoring

How do we choose blood thinners



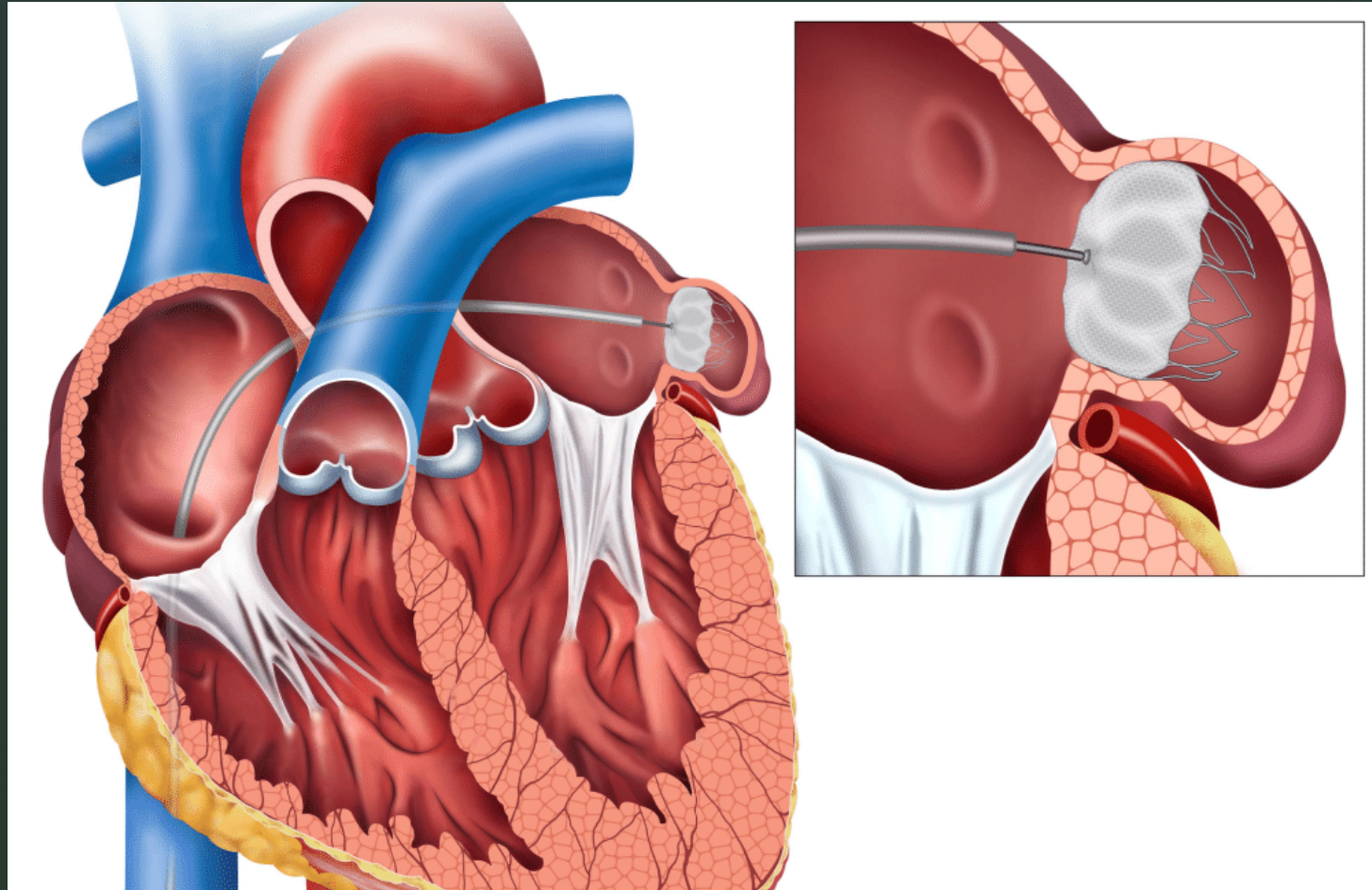
From Maan A, Heist EK, Ruskin JN, Mansour M. Practical issues in the management of novel oral anticoagulants – cardioversion and ablation. *J Thorac Dis* 2015;7:115–31, with permission from AME Publishing Company.

Key: AF = atrial fibrillation; FDA = Food and Drugs Administration; NOAC = non-vitamin K oral anticoagulant

Who should remain on warfarin?

- Patient already receiving warfarin and stable whose INR is easy to control
- If dabigatran, rivaroxaban, apixaban not available
- Cost
- If patient not likely to comply with twice daily dosing (Dabigatran, Apixaban)
- Chronic kidney disease (GFR < 30 ml/min)

WATCHMAN DEVICE: CLOSURE OF LEFT ATRIAL APPENDAGE





▶ RATE and RHYTHM CONTROL

Rate control as preferred therapy

- Age \geq 65, less symptomatic, hypertension
- Recurrent afib
- Previous antiarrhythmic drug failure
- Unlikely to maintain sinus rhythm (enlarged LA)

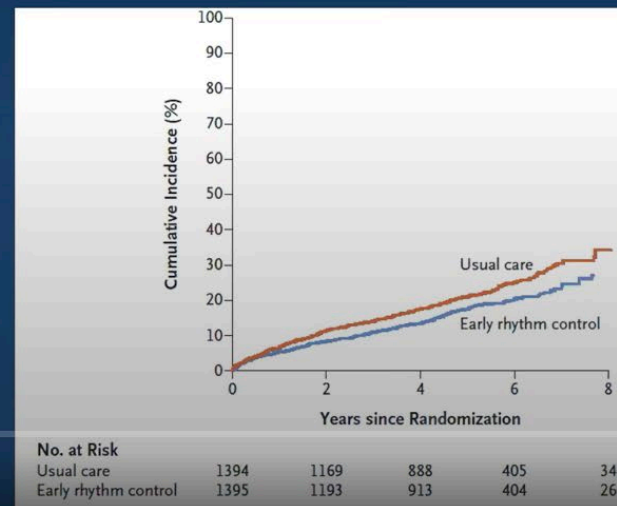
Rate Control Options

- Beta blocker
- Calcium channel blocker
 - Verapamil, diltiazem
- Digoxin
- AV junction ablation plus pacemaker

Early Rhythm-Control Therapy in Patients with Atrial Fibrillation

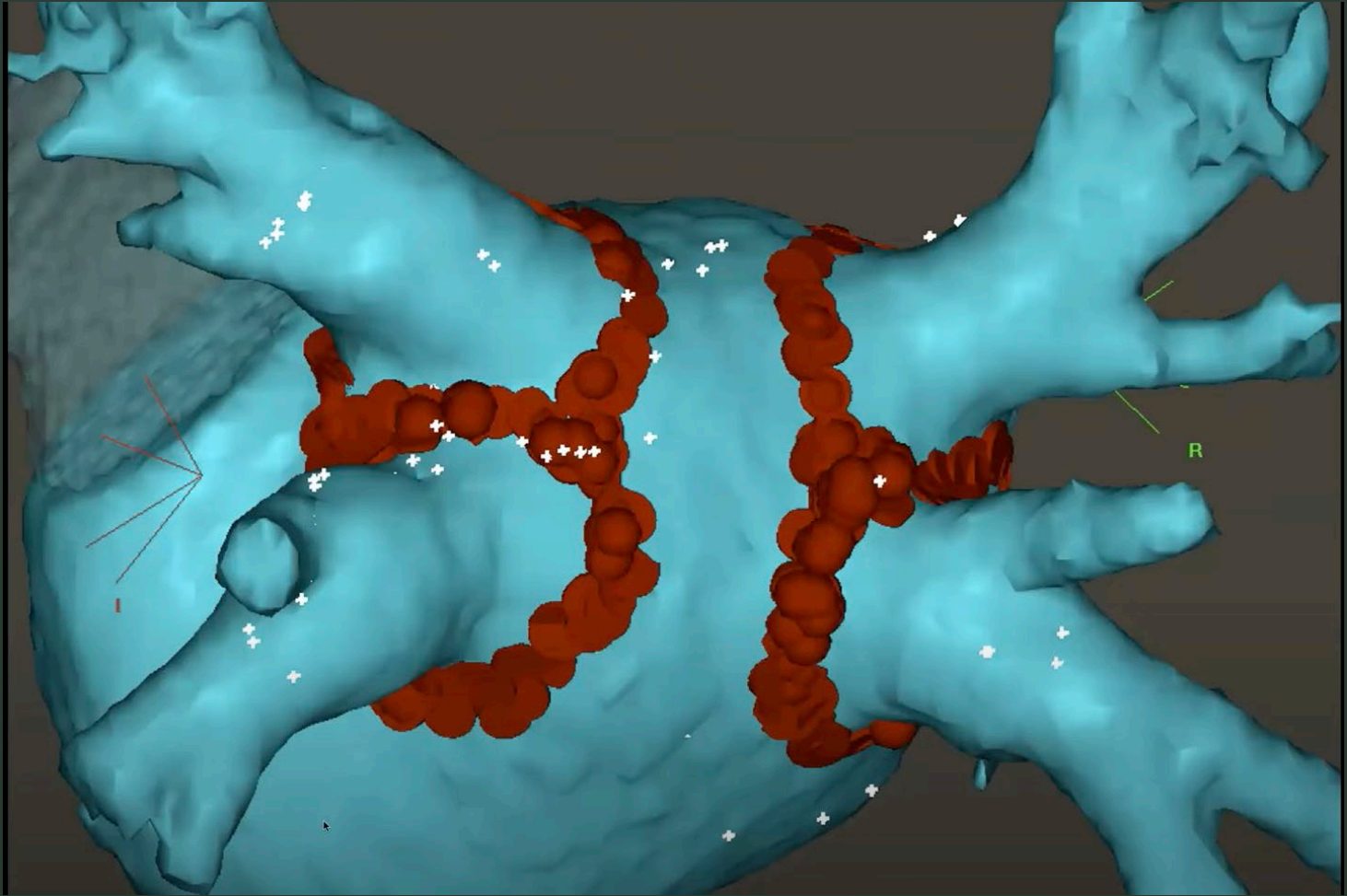
P. Kirchhof, A.J. Camm, A. Goette, A. Brandes, L. Eckardt, A. Elvan, T. Fetsch, I.C. van Gelder, D. Haase, L.M. Haegeli, F. Hamann, H. Heidbüchel, G. Hindricks, J. Kautzner, K.-H. Kuck, L. Mont, G.A. Ng, J. Rekosz, N. Schoen, U. Schotten, A. Suling, J. Taggeselle, S. Themistoclakis, E. Vettorazzi, P. Vardas, K. Wegscheider, S. Willems, H.J.G.M. Crijns, and G. Breithardt, for the EAST-AFNET 4 Trial Investigators*

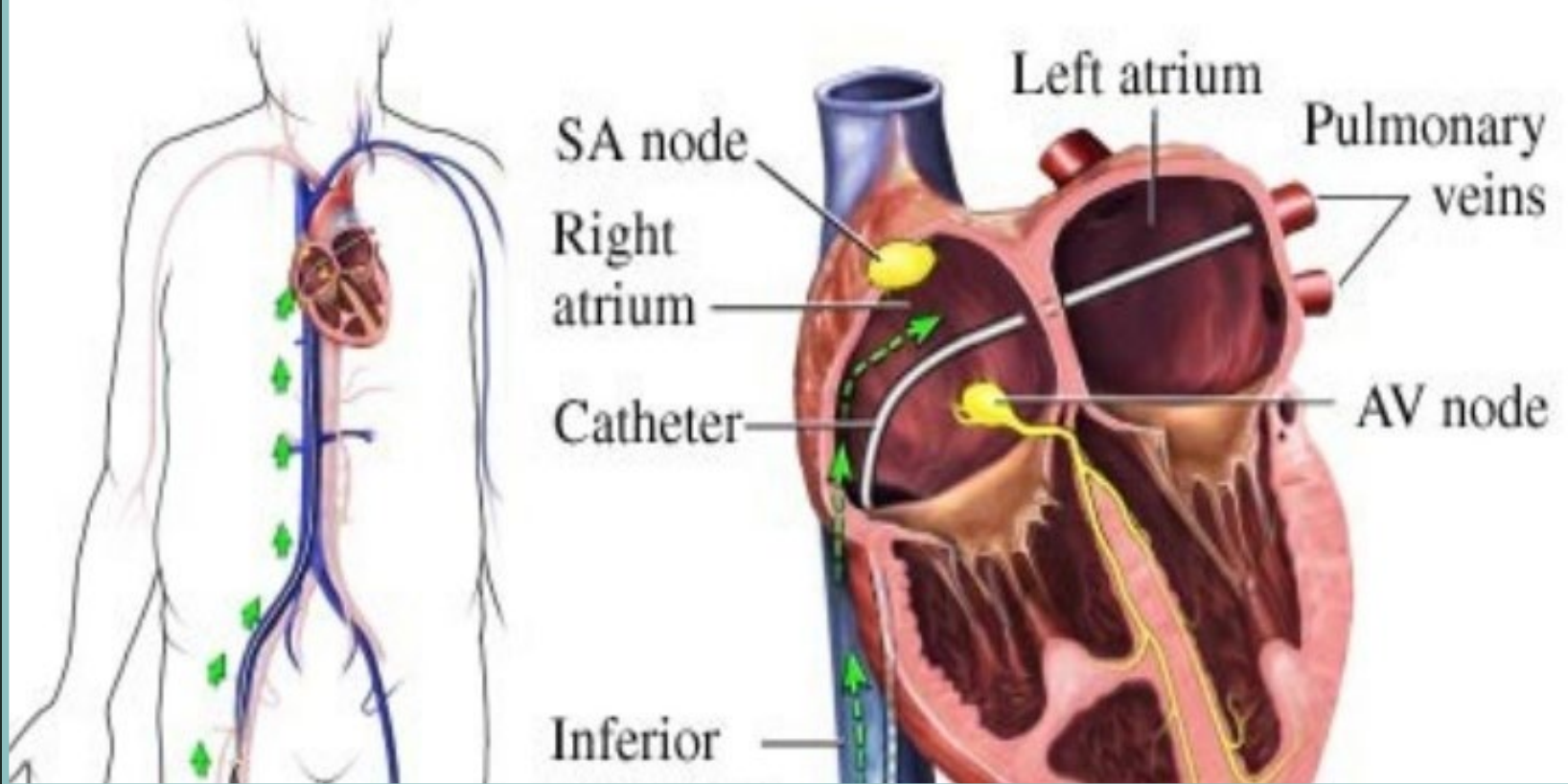
Primary outcome was a composite of CV death, stroke, heart failure hospitalization, or ACS
3.9 vs 5.0 per 100 pt years ($p < 0.01$)



Antiarrhythmic Medications

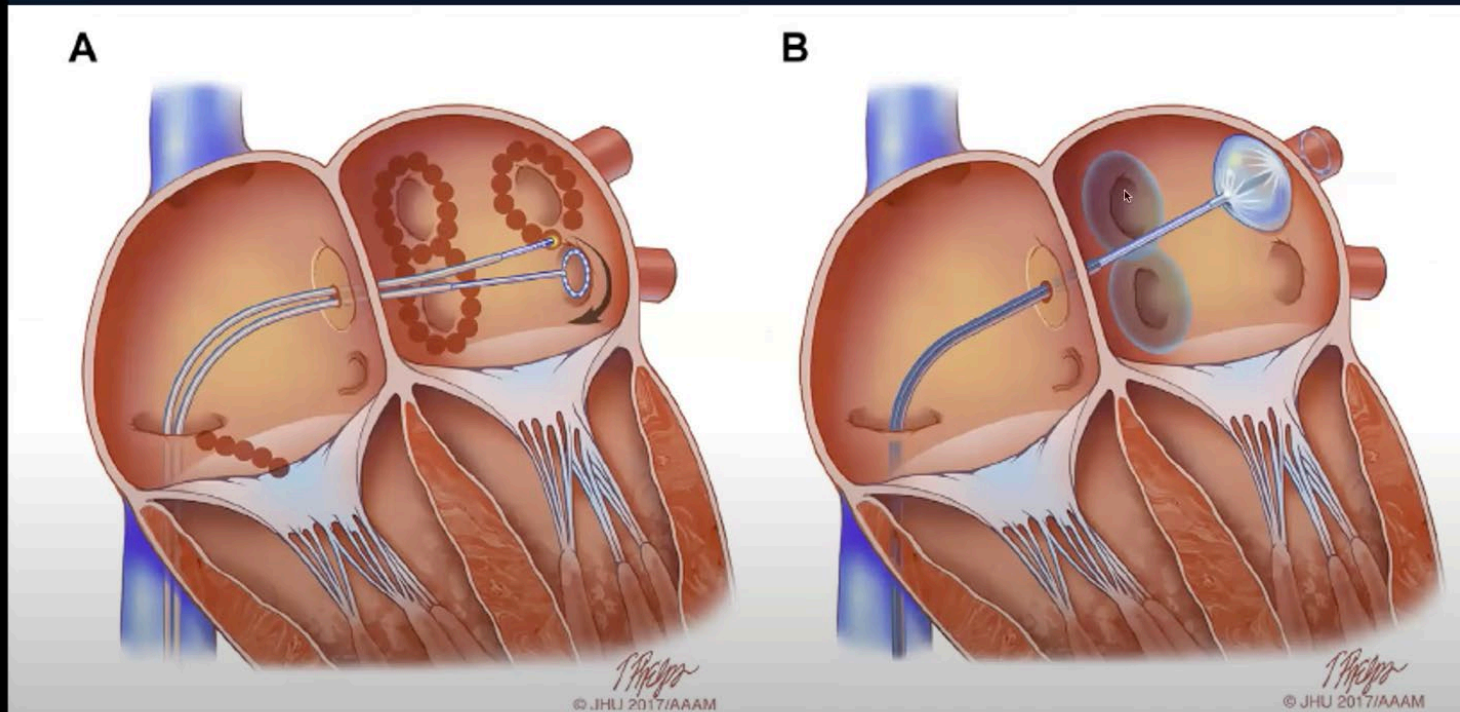
- Amiodarone
- Flecainide
- Multaq
- Propafenone
- Sotalol
- Tikosyn





A fib Ablation

ABLATION



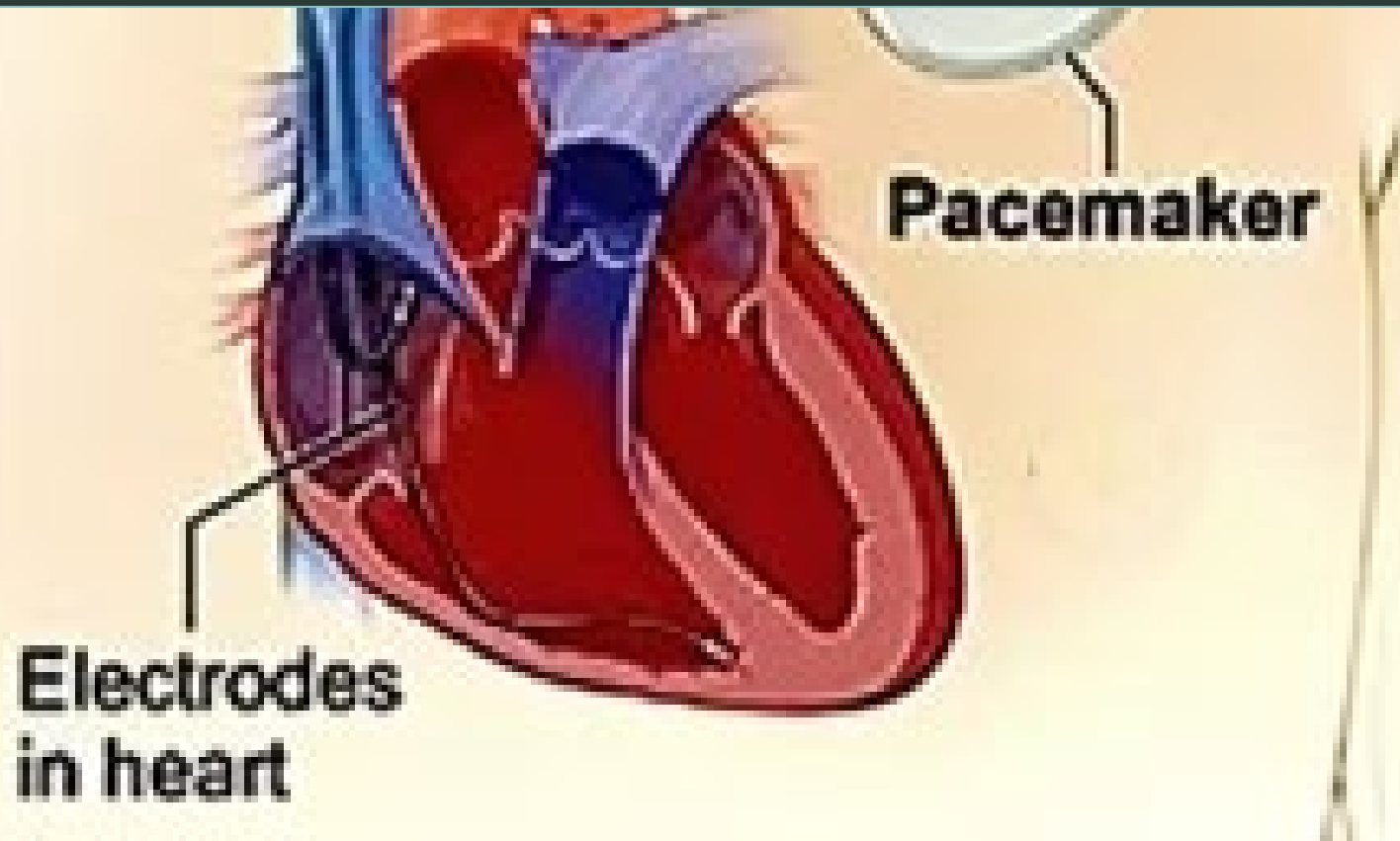
Treatment for permanent AF

- **Heart Rate control**

minimise symptoms associated with excessive heart rates

prevent tachycardia-associated cardiomyopathy

- **Anticoagulation**



Pacemaker with AV nodal Ablation



PREVENTION OF A FIB



LIFESTYLE FACTORS

- Obesity³²⁻³⁵
- Alcohol consumption^{3; 35; 36}
- Risks for cardiovascular disease: smoking, stress, caffeine and other stimulants³
- Activity level^{2; 3; 35}



OTHER CONDITIONS

- High blood pressure³⁵
- Heart failure^{27; 31; 37-40}
- History of heart attack^{27; 41}
- Coronary artery and other heart disease^{27; 33}
- Previous surgery^{42; 43}
- Sleep-disordered breathing (eg, obstructive sleep apnoea)^{35; 44}
- Diabetes^{35; 45}



NON-MODIFIABLE FACTORS

- Older age^{3; 46}
- Congenital heart defects⁴⁵
- Family history or other genetic factors^{27; 47; 48}
- Male sex^{3; 27; 46}



Thank You

